## **WEST Search History**

DATE: Friday, July 11, 2003

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	L11	L10 and (smooth muscle) and proliferat\$	12	L11
	L10	L8 and (polyurethane? or silicone? or acrylate? or polyester? or (polyalkylene oxide?) or polyalcohol? or polyolefin? or (polyvinyl chloride?) or cellulose? or polyamide? or polyolefin? or (fluorinated polymer?))	17	L10
	L9	L8 and biostable	. 1	L9
	L8	L7 and (polyhydroxy\$ or polyanhydride? or polyphosphazene? or polyphosphorester? or polyorthocarbonate? or polyamide? or polyorthocarbonate?)	17	L8
	L7	L6 and biodegrad\$	24	L7
	L6	L5 and (cerivastatin or atorvastatin or fluvastatin or lovastatin or pravastatin)	36	L6
	L5	L4 and (sol or gel)	47	L5
	L4	L3 and (polymer\$ or nonpolymer\$)	113	L4
	L3	L2 and coat\$	129	L3
	L2	L1 and stent	146	L2
	L1	(HMG-CoA) and inhibit\$	2918	L1

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=> s (HMG-COA) and inhibit?

L1 17156 (HMG-COA) AND INHIBIT?

=> s ll and stent

L2 183 L1 AND STENT

=> s 12 and coat?

L3 134 L2 AND COAT?

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L4 113 L3 AND (POLYMER OR NONPOLYMER)

=> s 14 and (sol or gel)

L5 41 L4 AND (SOL OR GEL)

=> s 15 and (cerivastatin or atorvastatin or fluvastatin or lovastatin or pravastatin)

L6 26 L5 AND (CERIVASTATIN OR ATORVASTATIN OR FLUVASTATIN OR LOVASTAT IN OR PRAVASTATIN)

=> s 16 and biodegrad

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17156 S (HMG-COA) AND INHIBIT?

L1 183 S L1 AND STENT

L2

L3 134 S L2 AND COAT?

113 S L3 AND (POLYMER OR NONPOLYMER) L4

L5 41 S L4 AND (SOL OR GEL)

26 S L5 AND (CERIVASTATIN OR ATORVASTATIN OR FLUVASTATIN OR LOVAS

=> s 16 and biodegrad?

18 L6 AND BIODEGRAD? L7

=> s 17 and (polyhydroxy? or polyanhydride# or polyphosphazene# or polyphophoester# or polyorthocarbonate# or polyamide# or polyorthoester# or polyphosphoester# or polyorthocarbonate#)

13 L7 AND (POLYHYDROXY? OR POLYANHYDRIDE# OR POLYPHOSPHAZENE# OR POLYPHOPHOESTER# OR POLYORTHOCARBONATE# OR POLYAMIDE# OR POLYORT HOESTER# OR POLYPHOSPHOESTER# OR POLYORTHOCARBONATE#)

=> s 18 and (polyurethane# or silicone# or acrylate# or polyestester# or (polyalkylene oxde#) or polyalcohol# or polyolefin# or (polyvinyl chloride#) or cellulose# or polyamide# or (fluorinated polymer#))

3 FILES SEARCHED...

13 L8 AND (POLYURETHANE# OR SILICONE# OR ACRYLATE# OR POLYESTESTER # OR (POLYALKYLENE OXDE#) OR POLYALCOHOL# OR POLYOLEFIN# OR (POLYVINYL CHLORIDE#) OR CELLULOSE# OR POLYAMIDE# OR (FLUORINATE D POLYMER#))

=> s 19 and ((smooth muscle) and proliferat?)

4 FILES SEARCHED...

9 L9 AND ((SMOOTH MUSCLE) AND PROLIFERAT?)

=> d 19 1-10 ibib abs

ANSWER 1 OF 13 USPATFULL

ACCESSION NUMBER: 2003:173192 USPATFULL

TITLE: Methods of using 48149, a human aminopeptidase family

INVENTOR(S): Chun, Miyoung, Belmont, MA, UNITED STATES

PATENT ASSIGNEE(S): Millennium Pharmaceuticals, Inc. (U.S. corporation)

NUMBER KIND DATE PATENT INFORMATION: US 2003119036 A1 20030626 APPLICATION INFO.: US 2002-281904 20021028 (10) A1

> NUMBER DATE \_\_\_\_\_\_

US 2001-335084P PRIORITY INFORMATION: 20011031 (60)

DOCUMENT TYPE: Utility FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: Steven A. Bossone, Millennium Pharmaceuticals, Inc., 75

Sidney Street, Cambridge, MA, 02139

NUMBER OF CLAIMS: 13 EXEMPLARY CLAIM: 1

7 Drawing Page(s) NUMBER OF DRAWINGS:

LINE COUNT:

5695

Isolated nucleic acids molecules, designated 48149 nucleic acid AB molecules, which encode human Aminopeptidase N, are disclosed. The invention provides methods of modulating 48149 activity, which is associated with the formation of atherosclerotic lesions in blood vessels. The invention further provides methods of treating, preventing and diagnosing cardiovascular disorders such as atherosclerosis, as well as disorders associated with the metabolism of lipids, for example, in the liver.

ANSWER 2 OF 13 USPATFULL

ACCESSION NUMBER:

2003:159431 USPATFULL

TITLE:

Methods of using 279, a human G protein-coupled protein

INVENTOR(S):

Logan, Thomas Joseph, Needham, MA, UNITED STATES

Galvin, Katherine M., Jamaica Plain, MA, UNITED STATES.

PATENT ASSIGNEE(S):

Millennium Pharmaceuticals, Inc. (U.S. corporation)

NUMBER KIND DATE \_\_\_\_\_\_ US 2003109044 20030612 Α1 US 2002-267811 20021009 (10) **A**1

PATENT INFORMATION: APPLICATION INFO .:

> NUMBER DATE

PRIORITY INFORMATION:

US 2001-329648P 20011016 (60)

DOCUMENT TYPE:

Utility

APPLICATION

FILE SEGMENT:

LEGAL REPRESENTATIVE:

MILLENNIUM PHARMACEUTICALS, INC., 75 Sidney Street,

Cambridge, MA, 02139

NUMBER OF CLAIMS:

EXEMPLARY CLAIM: NUMBER OF DRAWINGS:

7 Drawing Page(s)

LINE COUNT:

5366

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

The invention provides isolated nucleic acids molecules, designated 279 nucleic acid molecules, which encode human G protein-coupled receptor (GPCR) family members. The invention also provides antisense nucleic acid molecules, recombinant expression vectors containing 279 nucleic acid molecules, host cells into which the expression vectors have been introduced, and nonhuman transgenic animals in which a 279 gene has been introduced or disrupted. The invention still further provides isolated 279 proteins, fusion proteins, antigenic peptides and anti-279 antibodies. Methods utilizing compositions of the invention to treat, prevent or diagnose angiogenic disorders, e.g., cardiovascular and cancerous disorders, are also provided.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 3 OF 13 USPATFULL

ACCESSION NUMBER:

2003:140464 USPATFULL

TITLE:

Novel human membrane-associated protein and cell

surface protein family members

INVENTOR(S):

Meyers, Rachel E., Newton, MA, UNITED STATES

Glucksmann, Maria Alexandra, Lexington, MA, UNITED

STATES

Curtis, Rory A. J., Framingham, MA, UNITED STATES Kapeller-Libermann, Rosana, Chestnut Hill, MA, UNITED

STATES

Bandaru, Rajasekhar, Watertown, MA, UNITED STATES

Leiby, Kevin R., Natick, MA, UNITED STATES

NUMBER KIND DATE US 2003096305 A1 US 2002-162435 A1 PATENT INFORMATION: 20030522

APPLICATION INFO.: 20020604 (10)

US 2000-214220P

RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 2001-836499, filed

DATE

20000623 (60)

on 17 Apr 2001, PENDING

NUMBER

-----WO 2001-US12420 PRIORITY INFORMATION: 20010417

WO 2001-US19963 20010625 WO 2001-US16013 20010518 WO 2001-US20055 20010621

WO 2002-US275 20020108 WO 2001-US41811 20010821 US 2000-197507P 20000418 (60)

US 2000-205674P 20000519 (60) US 2000-213963P 20000623 (60) US 2001-260286P 20010108 (60)

US 2000-226612P 20000821 (60)

DOCUMENT TYPE: Utility FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: LOUIS MYERS, Fish & Richardson P.C., 225 Franklin

Street, Boston, MA, 02110-2804

NUMBER OF CLAIMS:

EXEMPLARY CLAIM:

NUMBER OF DRAWINGS: 22 Drawing Page(s)

LINE COUNT: 30445

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

The invention provides isolated nucleic acids molecules, designated 16051a, 16051b, 58199, 57805, 56739, 39362, and 23228 nucleic acid molecules, which encode novel human membrane-associated protein family members, and human cell surface protein family members. The invention also provides antisense nucleic acid molecules, recombinant expression vectors containing 1605la, 1605lb, 58199, 57805, 56739, 39362, or 23228 nucleic acid molecules, host cells into which the expression vectors have been introduced, and nonhuman transgenic animals in which a 1605la, 16051b, 58199, 57805, 56739, 39362, or 23228 gene has been introduced or disrupted. The invention still further provides isolated 16051a, 16051b, 58199, 57805, 56739, 39362, or 23228 proteins, fusion proteins; antigenic peptides and anti-16051a, 16051b, 58199, 57805, 56739, 39362, or 23228 antibodies. Diagnostic methods utilizing compositions of the invention are also provided.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 4 OF 13 USPATFULL

ACCESSION NUMBER: 2003:133526 USPATFULL

TITLE: Devices and compounds for treating arterial restenosis

INVENTOR(S): Zahradka, Peter, Winnipeg, CANADA

NUMBER KIND DATE \_\_\_\_\_\_\_ US 2003091611 A1 20030515 US 2002-273103 A1 20021016 (10) PATENT INFORMATION:

APPLICATION INFO.:

RELATED APPLN. INFO.: Division of Ser. No. US 2000-585886, filed on 31 May

2000, PENDING

NUMBER DATE -----

PRIORITY INFORMATION: US 1999-150696P 19990602 (60)

DOCUMENT TYPE:

Utility

FILE SEGMENT:

APPLICATION

LEGAL REPRESENTATIVE:

Nicholas S. Buffinger, Morrison & Foerster LLP, 755

Page Mill Road, Palo Alto, CA, 94304-1018

NUMBER OF CLAIMS:

EXEMPLARY CLAIM:

NUMBER OF DRAWINGS:

34 Drawing Page(s)

LINE COUNT:

2051

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

Described herein is the use of ADPRT decoy substrates to treat or prevent proliferative disorders. In one example, MIBG is shown to prevent restenosis in damaged vessels. In one embodiment, MIBG is combined with an adhesive agent for localizing the mixture to the site of injury. As a result of this arrangement, MIBG is not systemically released.

## CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 5 OF 13 USPATFULL

ACCESSION NUMBER:

2003:120257 USPATFULL

TITLE:

1983, 52881, 2398, 45449, 50289, and 52872, novel  ${\tt G}$ 

protein-coupled receptors and uses therefor

INVENTOR(S):

Glucksmann, Maria Alexandra, Lexington, MA, UNITED

STATES

Galvin, Katherine M., Jamaica Plain, MA, UNITED STATES

Silos-Santiago, Inmaculada, Cambridge, MA, UNITED

STATES

PATENT ASSIGNEE(S):

Millennium Pharmaceuticals, Inc., a Delaware

corporation (U.S. corporation)

NUMBER KIND DATE \_\_\_\_\_\_ US 2003082738 A1 20030501

PATENT INFORMATION:

US 2002-282837 A1 20021029 (10)

APPLICATION INFO.: RELATED APPLN. INFO.:

Continuation of Ser. No. US 2001-796338, filed on 28

Feb 2001, ABANDONED

NUMBER DATE \_\_\_\_\_\_

PRIORITY INFORMATION:

US 2000-186059P 20000229 (60)

DOCUMENT TYPE:

Utility

FILE SEGMENT:

APPLICATION

LEGAL REPRESENTATIVE:

FISH & RICHARDSON PC, 225 FRANKLIN ST, BOSTON, MA,

02110

NUMBER OF CLAIMS: EXEMPLARY CLAIM: 36 1

NUMBER OF DRAWINGS:

49 Drawing Page(s)

8096 LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

The invention provides isolated nucleic acids molecules, designated 1983, 52881, 2398, 45449, 50289, and 52872 nucleic acid molecules, which encode novel G protein-coupled receptor members. The invention also provides antisense nucleic acid molecules, recombinant expression vectors containing 1983, 52881, 2398, 45449, 50289, or 52872 nucleic acid molecules, host cells into which the expression vectors have been introduced, and nonhuman transgenic animals in which a 1983, 52881, 2398, 45449, 50289, or 52872 gene has been introduced or disrupted. The invention still further provides isolated 1983, 52881, 2398, 45449, 50289, or 52872 proteins, fusion proteins, antigenic peptides and anti-1983, 52881, 2398, 45449, 50289, or 52872 antibodies. Diagnostic methods utilizing compositions of the invention are also provided.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L9 ANSWER 6 OF 13 USPATFULL

ACCESSION NUMBER: 2003:112580 USPATFULL

TITLE: Stent coatings containing
HMG-CoA reductase inhibitors

INVENTOR(S): Pathak, Chandrashekhar, Austin, TX, UNITED STATES

Akella, Rama, Austin, TX, UNITED STATES Ranieri, John, Atlanta, GA, UNITED STATES

NUMBER KIND DATE

PATENT INFORMATION: US 2003077310 A1 20030424 APPLICATION INFO.: US 2001-991235 A1 20011022 (9)

DOCUMENT TYPE: Utility
FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: SULZER MEDICA USA INC., Suite 1600, 3 East Greenway

Plaza, Houston, TX, 77046

NUMBER OF CLAIMS: 41 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 7 Drawing Page(s)

LINE COUNT: 962

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Stents with coatings comprising a combination of a restenosis

inhibitor comprising an HMG-CoA reductase

inhibitor and a carrier. Also provided are methods of

coating stents with a combination of an HMG-

CoA reductase inhibitor and a carrier. A preferred example of a restenosis inhibitor is cerivastatin.

The stent coatings have been shown to release restenosis inhibitors in their active forms.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L9 ANSWER 7 OF 13 USPATFULL

ACCESSION NUMBER: 2003:95829 USPATFULL

TITLE: Devices and compounds for treating arterial restenosis

INVENTOR(S): Zahradka, Peter, Winnipeg, CANADA

PATENT ASSIGNEE(S): Cardiovascular Solutions, Inc., Winnipeg, CANADA

(non-U.S. corporation)

APPLICATION INFO.: US 2000-585886 20000531 (9)

NUMBER DATE

\_\_\_\_\_

PRIORITY INFORMATION: US 1999-150696P 19990602 (60)

DOCUMENT TYPE: Utility FILE SEGMENT: GRANTED

PRIMARY EXAMINER: Page, Thurman K.
ASSISTANT EXAMINER: Bennett, Rachel M.
LEGAL REPRESENTATIVE: Morrison & Foerster LLP

NUMBER OF CLAIMS: 6
EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 32 Drawing Figure(s); 34 Drawing Page(s)

LINE COUNT: 2271

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Described herein is the use of ADPRT decoy substrates to treat or prevent proliferative disorders. In one example, MIBG is shown to prevent restenosis in damaged vessels. In one embodiment, MIBG is

combined with an adhesive agent for localizing the mixture to the site of injury. As a result of this arrangement, MIBG is not systemically released.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L9 ANSWER 8 OF 13 USPATFULL

ACCESSION NUMBER: 2002:294662 USPATFULL

TITLE: 39362, a novel human CUB domain-containing protein

family member and uses thereof

INVENTOR(S): Bandaru, Rajasekhar, Watertown, MA, UNITED STATES

NUMBER DATE

PRIORITY INFORMATION: US 2001-260286P 20010108 (60)

DOCUMENT TYPE: Utility
FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: LOUIS MYERS, Fish & Richardson P.C., 225 Franklin

Street, Boston, MA, 02110-2804

NUMBER OF CLAIMS: 20 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 5 Drawing Page(s)

LINE COUNT: 5546

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

The invention provides isolated nucleic acids molecules, designated 39362 nucleic acid molecules, which encode novel CUB domain-containing protein members. The invention also provides antisense nucleic acid molecules, recombinant expression vectors containing 39362 nucleic acid molecules, host cells into which the expression vectors have been introduced, and nonhuman transgenic animals in which a 39362 gene has been introduced or disrupted. The invention still further provides isolated 39362 proteins, fusion proteins, antigenic peptides and anti-39362 antibodies. Diagnostic methods utilizing compositions of the invention are also provided.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L9 ANSWER 9 OF 13 USPATFULL

ACCESSION NUMBER: 2002:171640 USPATFULL

TITLE: Intravascular drug delivery device and use therefor

INVENTOR(S): Humes, H. David, Ann Arbor, MI, UNITED STATES

Tziampazis, Evangelos, Plymouth, MI, UNITED STATES

DOCUMENT TYPE: Utility
FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: TESTA, HURWITZ & THIBEAULT, LLP, HIGH STREET TOWER, 125

HIGH STREET, BOSTON, MA, 02110

NUMBER OF CLAIMS: 43 EXEMPLARY CLAIM: 1 NUMBER OF DRAWINGS: 9 Drawing Page(s)

LINE COUNT: 1905

Disclosed is an implantable drug delivery device for delivering a pre-selected drug directly into the systemic circulation of an animal. The device comprises an anchor immobilizable to an inner wall of an intact blood vessel. The device also comprises a drug containing reservoir that is retained in place within the blood vessel by the immobilized anchor. The reservoir may include, for example, a drug containing osmotic pump or a drug permeable capsule having disposed therein drug containing particles, which release the drug directly into blood passing the reservoir. The invention also provides a minimally invasive method for introducing into a blood vessel and, optionally, removing from the blood vessel the drug delivery device of the invention.

L9 ANSWER 10 OF 13 USPATFULL

ACCESSION NUMBER: 2002:157616 USPATFULL

TITLE: Lipid-based nitric oxide donors

INVENTOR(S): Herrmann, Robert A., Boston, MA, UNITED STATES

Naimark, Wendy, Cambridge, MA, UNITED STATES

PATENT ASSIGNEE(S): Scimed Life Systems, Inc. (U.S. corporation)

NUMBER KIND DATE ----------PATENT INFORMATION: US 2002082221 A1 20020627 APPLICATION INFO.: US 2000-745226 Α1 20001221 (9) DOCUMENT TYPE: Utility FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: Mayer, Fortkort & Williams, L.L.C., Suite 250, 200

Executive Drive, West Orange, NJ, 07052

NUMBER OF CLAIMS: 57 EXEMPLARY CLAIM: 1 LINE COUNT: 963

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Novel nitric-oxide releasing lipid molecules are provided which comprise a lipid molecule selected from (a) phosphoglycerides, (b) lipids having a sphingosine base as a backbone, (c) monoacylglyerols, (d) diacylglycerols, (e) glycosylacylglycerols, and (f) sterol compounds of the formula: ##STR1##

where R is a branched aliphatic chain of eight or more carbon atoms, wherein the lipid molecule is provided with a nitric-oxide containing group which comprises (a) a-S-N.dbd.O moiety, (b) a-O-N.dbd.O moiety, or (c) a ##STR2##

moiety. Also provided are methods of forming such nitric oxide releasing lipid molecules.

Various pharmaceutical compositions, topical liquids and drug delivery systems comprising the nitric-oxide releasing lipid molecules are also described. Further provided are methods for therapeutically administering nitric oxide to patients, methods of treating or preventing various conditions, methods for promoting wound healing and methods of reducing the cells present in an atherosclerotic lesion, which methods utilize the nitric-oxide releasing lipid molecules.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.